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visibility composition may further include additional radiopaque particles or contrast particles mixed in with the composition, which have a particle size between about 120µ and 350µ, preferably between ábout 120µ and 250µ.

## IN THE CLAIMS

Claims 33-43 should read as follows in view of this Amendment and the Second Preliminary Amendment filed December 17, 2001:

33. An injectable composition comprising:

a biocompatible matrix;

radiopaque particles mixed within said biocompatible matrix, said radiopaque particles having a particle size between about 120µ and 2200µ; and

liquid contrast agent,

- 34. The injectable composition of claim 33, wherein said biocompatible matrix and said radiopaque particles form a slurry.
- 35. The injectable composition of claim 33, wherein the mixture of said biocompatible matrix and said radiopaque particles forms a hard tissue implant material.
- 36. The injectable composition of claim 33, wherein said radiopaque particles have a particle size between about 350 µ and 2200 µ.
  - 37. The injectable composition of claim 36, further comprising:

radiopaque particles for contrast having a particle size between about 120µ and 350µ.

- 38. The injectable composition of claim 36, wherein said radiopaque particles have a particle size between about 450µ and 1600µ.
- 39. The injectable composition of claim 38, wherein said radiopaque particles having a particle size between about 570µ and 1150µ.
  - 40. An enhanced visibility composition comprising:
    - a flowable matrix; and

radiopaque particles in said flowable matrix, said radiopaque particles having a size between about 350µ and about 2200µ so as to be individually visible during implantation.

41. The enhanced visibility composition of claim 40, wherein said radiopaque particles have a size between about 570µ and 2200µ.

142. The enhanced visibility composition of claim 40, wherein said radiopaque particles have a size between about 450µ and 1600µ.



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New

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43. The enhanced visibility composition of claim 40, wherein said radiopaque particles have a size between about 570µ and 1150µ.

Please enter the following new claims, corresponding to original claims 33-41, except to the extent explained in the Remarks below

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744. (New) The injectable composition of claim 40, further comprising:

radiopaque particles for contrast having a particle size between about 120µ and 350µ.

45. (New) The injectable composition of claim 40, further comprising: radiopaque particles for contrast having a particle size up to about 350µ.

- 46. (New) The injectable composition of claim 36, further comprising: radiopaque particles for contrast having a particle size up to about 350µ.
- 47. (New) An injectable composition comprising:

a hard tissue implant biocompatible matrix; and

radiopaque particles mixed within said biocompatible matrix, said radiopaque particles having a particle size between about 120µ and 2200µ.

- 48. (New) The injectable composition of claim 47, wherein said biocompatible matrix and said radiopaque particles form a slurry.
- 49. (New) The injectable composition of claim 47, wherein said radiopaque particles have a particle size between about 350µ and 2200µ.
- 50. (New) The injectable composition of claim 47, wherein said radiopaque particles have a particle size between about 450μ and 1600μ.
- 51. (New) The injectable composition of claim 50, wherein said radiopaque particles have a particle size between about 570μ and 1150μ.
  - 52. (New) The injectable composition of claim 49, further comprising: radiopaque particles for contrast having a particle size between about 120μ and 350μ.
  - 53. (New) The injectable composition of claim 49, further comprising: radiopaque particles for contrast having a particle size up to about 350µ.

## REMARKS

## Formal Matters

Claims 33-53 are pending after entry of the amendments set forth herein.

Claims 33-41 were examined in the Office Action dated December 19,2001 prior to entry of the-Second Preliminary Amendment filed by Applicant on December 4, 2001. As such, claims 33-41 as